

Refine Search

Search Results -

Terms	Documents
L2 and (pelz or dinman or czaplinski).in.	5

Database: US Pre-Grant Publication Full-Text Database
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Search: L3

Recall Text
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Search History

DATE: Wednesday, July 28, 2004 [Printable Copy](#) [Create Case](#)

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side by side				result set
DB=PGPB,USPT,EPAB,JPAB,DWPI; PLUR=YES; OP=OR				
<u>L3</u>	L2 and (pelz or dinman or czaplinski).in.		5	<u>L3</u>
<u>L2</u>	L1 and (upf\$4 or nam7\$4 or sal1\$4 or ifs2\$4 or mof4\$4 or nmd2\$4 or isf1\$4 or sua1\$4 or sua6\$4)		40	<u>L2</u>
<u>L1</u>	(HELICAS\$4 OR MTT1\$4) AND (ERF\$4 OR (RELEAS\$4 same FACTO\$4))		500	<u>L1</u>

END OF SEARCH HISTORY

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Search Results - Record(s) 1 through 5 of 5 returned.

1. Document ID: US 20040115787 A1

Using default format because multiple data bases are involved.

L3: Entry 1 of 5

File: PGPB

Jun 17, 2004

PGPUB-DOCUMENT-NUMBER: 20040115787

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040115787 A1

TITLE: Subfamily of RNA helicases which are modulators of the fidelity of translation termination and uses thereof

PUBLICATION-DATE: June 17, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peltz, Stuart	Piscataway	NJ	US	
<u>Czaplinski</u> , Kevin	Somerset	NJ	US	
Dinman, Jonathan D.	North Brunswick	NJ	US	

US-CL-CURRENT: 435/226; 530/388.26, 536/23.2

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	RIMC	Drawn D.
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2. Document ID: US 20030032158 A1

L3: Entry 2 of 5

File: PGPB

Feb 13, 2003

PGPUB-DOCUMENT-NUMBER: 20030032158

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030032158 A1

TITLE: Method of modulating the efficiency of translation termination and degradation of aberrant mRNA involving a surveillance complex comprising human Upf1p, eucaryotic release factor 1 and eucaryotic release factor 3

PUBLICATION-DATE: February 13, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peltz, Stuart	Piscataway	NJ	US	
<u>Czaplinski</u> , Kevin	Somerset	NJ	US	

Weng, Youmin Cranford NJ US

US-CL-CURRENT: 435/189; 530/388.26

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KOMC](#) | [Draw. D.](#)

3. Document ID: US 6630294 B1

L3: Entry 3 of 5

File: USPT

Oct 7, 2003

US-PAT-NO: 6630294

DOCUMENT-IDENTIFIER: US 6630294 B1

TITLE: Subfamily of RNA helicases which are modulators of the fidelity of translation termination and uses thereof

DATE-ISSUED: October 7, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Peltz; Stuart	Piscataway	NJ		
<u>Czaplinski</u> ; Kevin	Somerset	NJ		
<u>Dinman</u> ; Jonathan D.	North Brunswick	NJ		

US-CL-CURRENT: 435/4; 435/183, 435/7.1, 435/7.31, 436/86, 530/350, 536/23.2

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KOMC](#) | [Draw. D.](#)

4. Document ID: US 6486305 B1

L3: Entry 4 of 5

File: USPT

Nov 26, 2002

US-PAT-NO: 6486305

DOCUMENT-IDENTIFIER: US 6486305 B1

TITLE: METHOD OF MODULATING THE EFFICIENCY OF TRANSLATION TERMINATION AND DEGRADATION OF ABERRANT mRNA INVOLVING A SURVEILLANCE COMPLEX COMPRISING HUMAN UPF1P, EUKARYOTIC RELEASE FACTOR 1 AND EUKARYOTIC RELEASE FACTOR 3

DATE-ISSUED: November 26, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Peltz; Stuart	Piscataway	NJ	08854	
<u>Czaplinski</u> ; Kevin	Somerset	NJ	08873	
Weng; Youmin	Cranford	NJ	07016	

US-CL-CURRENT: 530/412; 435/455, 435/69.1, 530/350, 530/358

[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sequences](#) | [Attachments](#) | [Claims](#) | [KOMC](#) | [Draw. D.](#)

5. Document ID: US 20040115787 A1

L3: Entry 5 of 5

File: DWPI

Jun 17, 2004

DERWENT-ACC-NO: 2004-449400

DERWENT-WEEK: 200442

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TITLE: Identifying a test composition or agent that modulates the efficiency of translation termination comprises contacting the MTT1 with the test composition or agent, and determining if the test composition or agent inhibits the MTT1INVENTOR: CZAPLINSKI, K; DINMAN, J D ; PELTZ, SPRIORITY-DATA: 1998US-093685P (July 22, 1998), 1999US-0359268 (July 22, 1999),
2003US-0652334 (August 28, 2003)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
<u>US 20040115787 A1</u>	June 17, 2004		041	C12N009/64

INT-CL (IPC): C07 H 21/04; C07 K 16/40; C12 N 9/64[Full](#) | [Title](#) | [Citation](#) | [Front](#) | [Review](#) | [Classification](#) | [Date](#) | [Reference](#) | [Sentences](#) | [Attachments](#) | [Claims](#) | [KINDC](#) | [Drawn D.](#)[Clear](#)[Generate Collection](#)[Print](#)[Fwd Refs](#)[Bkwd Refs](#)[Generate OACS](#)

Terms

Documents

L2 and (pelz or dinman or czaplinski).in.

5

Display Format: [Change Format](#)[Previous Page](#)[Next Page](#)[Go to Doc#](#)

=> d his

(FILE 'HOME' ENTERED AT 19:31:12 ON 28 JUL 2004)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI,
BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA,
CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS,
DDFB, DDFU, DGENE, DRUGB, DRUGMONOG2, ...' ENTERED AT 19:31:26 ON 28 JUL
2004

SEA (HELICAS? OR MTT1?) OR (ERF? OR (RELEAS?(S)FACTO?))

1596 FILE ADISCTI
354 FILE ADISINSIGHT
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44363 FILE PASCAL
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475 FILE RDISCLOSURE
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4638 FILE WPIDS
1925 FILE WPIFV
4638 FILE WPINDEX
944 FILE IPA
171 FILE NAPRALERT
19988 FILE NLDB
L1 QUE (HELICAS? OR MTT1?) OR (ERF? OR (RELEAS?(S) FACTO?))

FILE 'DGENE, PROMT, EMBASE, CEABA-VTB, MEDLINE, SCISEARCH, CAPLUS,
PASCAL, BIOSIS, USPATFULL, ESBIOBASE' ENTERED AT 19:35:08 ON 28 JUL 2004
L2 673 S (HELICAS? OR MTT1?) AND (ERF? OR (RELEAS?(S)FACTO?))
L3 51 S L2 AND (UPF? OR NAM7? OR SAL1? OR IFS2? OR MOF4? OR NMD2? OR
L4 46 DUP REM L3 (5 DUPLICATES REMOVED)

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| NEWS | 5 | May 27 New UPM (Update Code Maximum) field for more efficient patent SDIs in Cplus |
| NEWS | 6 | May 27 Cplus super roles and document types searchable in REGISTRY |
| NEWS | 7 | Jun 28 Additional enzyme-catalyzed reactions added to CASREACT |
| NEWS | 8 | Jun 28 ANTE, AQUALINE, BIOENG, CIVILENG, ENVIROENG, MECHENG, and WATER from CSA now available on STN(R) |
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| NEWS EXPRESS | | MARCH 31 CURRENT WINDOWS VERSION IS V7.00A, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0JC(JP), AND CURRENT DISCOVER FILE IS DATED 26 APRIL 2004 |
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=> index bioscience medicine
FILE 'DRUGMONOG' ACCESS NOT AUTHORIZED
COST IN U.S. DOLLARS

FULL ESTIMATED COST

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU, DGENE, DRUGB, DRUGMONOG2, ...' ENTERED AT 19:31:26 ON 28 JUL 2004

73 FILES IN THE FILE LIST IN STNINDEX

Enter SET DETAIL ON to see search term postings or to view search error messages that display as 0* with SET DETAIL OFF.

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=> s (helicas? or mttl1?) or (erf? or (releas?(s)facto?))  
    1596 FILE ADISCTI  
    354 FILE ADISINSIGHT  
    378 FILE ADISNEWS  
   6851 FILE AGRICOLA  
   192 FILE ANABSTR  
  2954 FILE AQUASCI
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1469 FILE BIOBUSINESS
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35 FILES SEARCHED...
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L1 QUE (HELICAS? OR MTT1?) OR (ERF? OR (RELEAS?(S) FACTO?))

=> d rank

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|-----|--------|-------------|
| F1 | 399612 | DGENE |
| F2 | 304170 | PROMT |
| F3 | 83157 | EMBASE |
| F4 | 60171 | CEABA-VTB |
| F5 | 59476 | MEDLINE |
| F6 | 58511 | SCISEARCH |
| F7 | 50487 | CAPLUS |
| F8 | 44363 | PASCAL |
| F9 | 43311 | BIOSIS |
| F10 | 43056 | USPATFULL |
| F11 | 27817 | ESBIOBASE |
| F12 | 25707 | BIOTECHNO |
| F13 | 22758 | CANCERLIT |
| F14 | 21744 | DRUGU |
| F15 | 20977 | LIFESCI |
| F16 | 20905 | TOXCENTER |
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| F18 | 18653 | DDFU |
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| F20 | 14587 | CABA |
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| F22 | 10295 | DRUGB |
| F23 | 6851 | AGRICOLA |
| F24 | 4638 | WPIDS |
| F25 | 4638 | WPINDEX |
| F26 | 4438 | DISSABS |
| F27 | 4377 | JICST-EPLUS |
| F28 | 3955* | FEDRIP |
| F29 | 3489 | NTIS |
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| F48 | 475 | FROSTI |
| F49 | 475 | RDISCLOSURE |
| F50 | 455 | CROPU |
| F51 | 453 | HEALSAFE |
| F52 | 444 | CIN |
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| F54 | 378 | ADISNEWS |
| F55 | 354 | ADISINSIGHT |
| F56 | 306 | CROPB |
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| F59 | 234 | BIOCOMMERCE |
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| F61 | 192 | IMSRESEARCH |
| F62 | 183 | IMSDRUGNEWS |
| F63 | 171 | NAPRALERT |
| F64 | 130 | PHARMAML |
| F65 | 83 | CEN |
| F66 | 76 | MEDICONF |
| F67 | 72 | IMSPRODUCT |
| F68 | 41 | DRUGMONOG2 |

F69 11 NUTRACEUT
F70 10 SYNTHLINE
F71 5 PHIC

=> file f1-f11
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FULL ESTIMATED COST ENTRY SESSION
3.42 3.63

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8 FILES SEARCHED...
L2 673 (HELCAS? OR MTT1?) AND (ERF? OR (RELEAS?(S) FACTO?))

=> s l2 and (upf? or nam7? or sal1? or ifs2? or mof4? or nmd2? or isf1? or sua1? or sua6?)
L3 51 L2 AND (UPF? OR NAM7? OR SAL1? OR IFS2? OR MOF4? OR NMD2? OR
ISF1? OR SUA1? OR SUA6?)

=> dup rem 13
DUPLICATE IS NOT AVAILABLE IN 'DGENE'.
ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE
PROCESSING COMPLETED FOR L3
L4 46 DUP REM L3 (5 DUPLICATES REMOVED)

=> d ti l4 1-46

L4 ANSWER 1 OF 46 USPATFULL on STN
TI Methods for in vitro expansion and transdifferentiation of human
pancreatic acinar cells into insulin-producing cells

L4 ANSWER 2 OF 46 USPATFULL on STN
TI Ncc2705-the genome of a bifidobacterium

L4 ANSWER 3 OF 46 USPATFULL on STN

TI Subfamily of RNA **helicases** which are modulators of the fidelity of translation termination and uses thereof

L4 ANSWER 4 OF 46 USPATFULL on STN
TI Targets for therapeutic intervention identified in the mitochondrial proteome

L4 ANSWER 5 OF 46 USPATFULL on STN
TI Whole cell engineering by mutagenizing a substantial portion of a starting genome, combining mutations, and optionally repeating

L4 ANSWER 6 OF 46 USPATFULL on STN
TI Methods of identifying compounds that inhibit nonstop degradation of mRNA

L4 ANSWER 7 OF 46 USPATFULL on STN
TI Wound healing biomarkers

L4 ANSWER 8 OF 46 USPATFULL on STN
TI Methods of diagnosis of breast cancer, compositions and methods of screening for modulators of breast cancer

L4 ANSWER 9 OF 46 USPATFULL on STN
TI Composition for the detection of signaling pathway gene expression

L4 ANSWER 10 OF 46 USPATFULL on STN
TI Novel human polynucleotides and polypeptides encoded thereby

L4 ANSWER 11 OF 46 USPATFULL on STN
TI Methods of diagnosis of ovarian cancer, compositions and methods of screening for modulators of ovarian cancer

L4 ANSWER 12 OF 46 USPATFULL on STN
TI Novel full-length cDNA

L4 ANSWER 13 OF 46 USPATFULL on STN
TI Nucleic acid sequences relating to Candida albicans for diagnostics and therapeutics

L4 ANSWER 14 OF 46 USPATFULL on STN
TI Nucleic acid molecule and encoded protein associated with sterol synthesis and metabolism

L4 ANSWER 15 OF 46 USPATFULL on STN
TI DNA array sequence selection

L4 ANSWER 16 OF 46 MEDLINE on STN
TI Leaky termination at premature stop codons antagonizes nonsense-mediated mRNA decay in S. cerevisiae.

L4 ANSWER 17 OF 46 USPATFULL on STN
TI Novel full length cDNA

L4 ANSWER 18 OF 46 USPATFULL on STN
TI Novel methods of diagnosis of metastatic colorectal cancer, compositions and methods of screening for modulators of metastatic colorectal cancer

L4 ANSWER 19 OF 46 USPATFULL on STN
TI Protein-protein interactions in adipocyte cells (3)

L4 ANSWER 20 OF 46 USPATFULL on STN
TI Novel full-length cDNA

L4 ANSWER 21 OF 46 USPATFULL on STN
TI Segments of the human gene for telomerase reverse transcriptase

L4 ANSWER 22 OF 46 USPATFULL on STN
TI Yeast proteome analysis

L4 ANSWER 23 OF 46 USPATFULL on STN
TI Novel nucleic acids and polypeptides

L4 ANSWER 24 OF 46 USPATFULL on STN
TI Libraries of expressible gene sequences

L4 ANSWER 25 OF 46 USPATFULL on STN
TI Methods of diagnosis of ovarian cancer, compositions and methods of screening for modulators of ovarian cancer

L4 ANSWER 26 OF 46 USPATFULL on STN
TI Libraries of expressible gene sequences

L4 ANSWER 27 OF 46 USPATFULL on STN
TI Human genes and gene expression products

L4 ANSWER 28 OF 46 USPATFULL on STN
TI Protein-protein interactions in adipocyte cells

L4 ANSWER 29 OF 46 USPATFULL on STN
TI Method of modulating the efficiency of translation termination and degradation of aberrant mRNA involving a surveillance complex comprising human **Upf1p, eucaryotic release factor 1** and **eucaryotic release factor 3**

L4 ANSWER 30 OF 46 USPATFULL on STN
TI Subfamily of RNA **helicases** which are modulators of the fidelity of translation termination and uses thereof

L4 ANSWER 31 OF 46 USPATFULL on STN
TI Nucleic acid and amino acid sequences relating to *Enterococcus faecalis* for diagnostics and therapeutics

L4 ANSWER 32 OF 46 USPATFULL on STN
TI Cells immortalized with telomerase reverse transcriptase for use in drug screening

L4 ANSWER 33 OF 46 USPATFULL on STN
TI Promoter for telomerase reverse transcriptase

L4 ANSWER 34 OF 46 USPATFULL on STN
TI *ENTEROCOCCUS FAECALIS* POLYNUCLEOTIDES AND POLYPEPTIDES

L4 ANSWER 35 OF 46 USPATFULL on STN
TI Composition for the detection of signaling pathway gene expression

L4 ANSWER 36 OF 46 USPATFULL on STN
TI METHOD OF MODULATING THE EFFICIENCY OF TRANSLATION TERMINATION AND DEGRADATION OF ABERRANT MRNA INVOLVING A SURVEILLANCE COMPLEX COMPRISING HUMAN **UPF1P, EUCARYOTIC RELEASE FACTOR 1** AND **EUCARYOTIC RELEASE FACTOR 3**

L4 ANSWER 37 OF 46 USPATFULL on STN
TI Polynucleotides and polypeptides derived from corn ear

L4 ANSWER 38 OF 46 USPATFULL on STN
TI Genomic DNA sequences of *ashbya gossypii* and uses thereof

L4 ANSWER 39 OF 46 CAPLUS COPYRIGHT 2004 ACS on STN
TI Subfamily of RNA **helicases** which are modulators of the fidelity of translation termination

L4 ANSWER 40 OF 46 USPATFULL on STN
TI Telomerase catalytic subunit

L4 ANSWER 41 OF 46 MEDLINE on STN DUPLICATE 1
TI **Mtt1** is a **Upf1-like helicase** that interacts with the translation termination factors and whose overexpression can modulate termination efficiency.

L4 ANSWER 42 OF 46 MEDLINE on STN
TI RNA surveillance. Unforeseen consequences for gene expression, inherited genetic disorders and cancer.

L4 ANSWER 43 OF 46 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.
on STN DUPLICATE 2

TI A mutated human homologue to yeast **Upf1** protein has a dominant-negative effect on the decay of nonsense-containing mRNAs in mammalian cells.

L4 ANSWER 44 OF 46 MEDLINE on STN
TI The surveillance complex interacts with the translation **release factors** to enhance termination and degrade aberrant mRNAs.

L4 ANSWER 45 OF 46 SCISEARCH COPYRIGHT 2004 THOMSON ISI on STN
TI PURIFICATION AND CHARACTERIZATION OF THE **UPF1** PROTEIN - A FACTOR INVOLVED IN TRANSLATION AND MESSENGER-RNA DEGRADATION

L4 ANSWER 46 OF 46 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
TI New multiprotein complex which can modulate peptidyl transferase activity during translation, useful to treat diseases associated with peptidyl transferase activity e.g. Duchene/Becker Muscular Dystrophy -

=> d ibib abs 14 29 30 36 39 43 44 46

L4 ANSWER 29 OF 46 USPATFULL on STN
ACCESSION NUMBER: 2003:44848 USPATFULL
TITLE: Method of modulating the efficiency of translation termination and degradation of aberrant mRNA involving a surveillance complex comprising human **Upf1p**, eucaryotic release factor 1 and eucaryotic release factor 3

INVENTOR(S): Peltz, Stuart, Piscataway, NJ, UNITED STATES
Czaplinski, Kevin, Somerset, NJ, UNITED STATES

PATENT ASSIGNEE(S): Weng, Youmin, Cranford, NJ, UNITED STATES
University of Medicine and Dentistry of New Jersey, New Brunswick, NY, UNITED STATES, 08903 (U.S. corporation)

| | NUMBER | KIND | DATE |
|-----------------------|--|------|---------------|
| PATENT INFORMATION: | US 2003032158 | A1 | 20030213 |
| APPLICATION INFO.: | US 2002-138784 | A1 | 20020503 (10) |
| RELATED APPLN. INFO.: | Continuation of Ser. No. US 1999-321649, filed on 28 May 1999, ABANDONED | | |

| | NUMBER | DATE |
|-----------------------|---|---------------|
| PRIORITY INFORMATION: | US 1998-86986P | 19980528 (60) |
| DOCUMENT TYPE: | Utility | |
| FILE SEGMENT: | APPLICATION | |
| LEGAL REPRESENTATIVE: | PERKINS COIE LLP, POST OFFICE BOX 1208, SEATTLE, WA, 98111-1208 | |

| | |
|---------------------|--------------------|
| NUMBER OF CLAIMS: | 28 |
| EXEMPLARY CLAIM: | 1 |
| NUMBER OF DRAWINGS: | 11 Drawing Page(s) |
| LINE COUNT: | 2935 |

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Provided are novel methods and assays to identify agents and compositions that modulate the ability of the eukaryotic surveillance complex to effect translation termination and degradation of aberrant mRNA.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L4 ANSWER 30 OF 46 USPATFULL on STN
ACCESSION NUMBER: 2003:268126 USPATFULL
TITLE: Subfamily of RNA **helicases** which are modulators of the fidelity of translation termination and uses thereof

INVENTOR(S): Peltz, Stuart, Piscataway, NJ, United States
Czaplinski, Kevin, Somerset, NJ, United States

PATENT ASSIGNEE(S): Dinman, Jonathan D., North Brunswick, NJ, United States
University of Medicine and Dentistry of New Jersey, New

Brunswick, NJ, United States (U.S. corporation)

| | NUMBER | KIND | DATE |
|---------------------|----------------|------|--------------|
| PATENT INFORMATION: | US 6630294 | B1 | 20031007 |
| APPLICATION INFO.: | US 1999-359268 | | 19990722 (9) |

| | NUMBER | DATE |
|-----------------------|--|---------------|
| PRIORITY INFORMATION: | US 1998-93685P | 19980722 (60) |
| DOCUMENT TYPE: | Utility | |
| FILE SEGMENT: | GRANTED | |
| PRIMARY EXAMINER: | Prouty, Rebecca E. | |
| ASSISTANT EXAMINER: | Ramirez, Delia | |
| LEGAL REPRESENTATIVE: | Wise, Michael J., Perkins Coie LLP | |
| NUMBER OF CLAIMS: | 4 | |
| EXEMPLARY CLAIM: | 1 | |
| NUMBER OF DRAWINGS: | 6 Drawing Figure(s); 6 Drawing Page(s) | |
| LINE COUNT: | 2768 | |

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention provides a method for modulating the efficiency of translation termination of messenger RNA. Also provided are methods of screening for compositions and agents capable of modulating translation termination.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L4 ANSWER 36 OF 46 USPATFULL on STN
ACCESSION NUMBER: 2002:311029 USPATFULL
TITLE: METHOD OF MODULATING THE EFFICIENCY OF TRANSLATION TERMINATION AND DEGRADATION OF ABERRANT mRNA INVOLVING A SURVEILLANCE COMPLEX COMPRISING HUMAN UPF1P , EUCLYTIC RELEASE FACTOR 1 AND EUCLYTIC RELEASE FACTOR 3
INVENTOR(S): Peltz, Stuart, 67 Castle Pointe Blvd., Piscataway, NJ, United States 08854
Czapinski, Kevin, 115 Hollywood Ave., Somerset, NJ, United States 08873
Weng, Youmin, 2 Indian Spring Rd., Cranford, NJ, United States 07016

| | NUMBER | KIND | DATE |
|-----------------------|---|------|--------------|
| PATENT INFORMATION: | US 6486305 | B1 | 20021126 |
| APPLICATION INFO.: | US 2000-639987 | | 20000816 (9) |
| RELATED APPLN. INFO.: | Division of Ser. No. US 1998-86260, filed on 28 May 1998, now abandoned | | |
| DOCUMENT TYPE: | Utility | | |
| FILE SEGMENT: | GRANTED | | |
| PRIMARY EXAMINER: | McCarry, Sean | | |
| ASSISTANT EXAMINER: | Zara, Jane | | |
| LEGAL REPRESENTATIVE: | Lyon & Lyon LLP | | |
| NUMBER OF CLAIMS: | 3 | | |
| EXEMPLARY CLAIM: | 1 | | |
| NUMBER OF DRAWINGS: | 11 Drawing Figure(s); 11 Drawing Page(s) | | |
| LINE COUNT: | 2808 | | |

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention provides a method of modulating translation termination efficiency of mRNA and/or promoting degradation of aberrant transcripts. Also, this invention provides a method of screening for a drug active involved in enhancing translation termination and a method for identifying a disease state involving defective the protein complex.

This invention provides a purified complex comprising an amount of a human Upf1p protein, a peptidyl eucaryotic release factor 1 (eRF1) and a peptidyl eucaryotic release factor 3 (eRF3) effective to modulate translation termination. Further, this invention provides an expression vector which comprises a nucleic acid encoding a human Upf1p protein, a peptidyl eucaryotic release factor 1 (eRF1) and a peptidyl eucaryotic

release factor 3 (eRF3) operably linked to a regulatory element.

This invention provides an antibody which binds to the complex comprising an amount of a human **Upf1p** protein, a peptidyl eucaryotic **release factor 1 (eRF1)** and a peptidyl eucaryotic **release factor 3 (eRF3)**) effective to modulate translation termination. This invention provides an agent which inhibits or modulates the binding of human **Upf1p** to **eRF1** or **eRF3** The agent may inhibit or facilitate the binding of human **Upf1p** to **eRF1** or **eRF3**

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L4 ANSWER 39 OF 46 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2000:85103 CAPLUS
DOCUMENT NUMBER: 132:148498
TITLE: Subfamily of RNA **helicases** which are modulators of the fidelity of translation termination
INVENTOR(S): Peltz, Stuart; Czaplinski, Kevin; Dinman, Jonathan D.
PATENT ASSIGNEE(S): University of Medicine and Dentistry, USA
SOURCE: PCT Int. Appl., 89 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|-------------|
| WO 2000005586 | A2 | 20000203 | WO 1999-US16802 | 19990722 |
| WO 2000005586 | A3 | 20000420 | | |
| W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | | | | |
| RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | | | | |
| CA 2338312 | AA | 20000203 | CA 1999-2338312 | 19990722 |
| AU 9952286 | A1 | 20000214 | AU 1999-52286 | 19990722 |
| EP 1098905 | A2 | 20010516 | EP 1999-937450 | 19990722 |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO | | | | |
| JP 2002524719 | T2 | 20020806 | JP 2000-561501 | 19990722 |
| PRIORITY APPLN. INFO.: | | | US 1998-120435 | A2 19980722 |
| | | | WO 1999-US16802 | W 19990722 |

AB This invention provides a method of modulating translation termination efficiency of mRNA and/or promoting degrdn. of aberrant transcripts. Also, this invention provides a method of screening for a drug active involved in enhancing translation termination and a method for identifying a disease state involving defective the protein complex. This invention provides a purified complex comprising an amt. of **MTT1** (mediator of translation termination, the gene encoding **helicase B**), human **Upf1p**, a peptidyl eukaryotic **release factor 1 (eRF1)** and a peptidyl eukaryotic **release factor 3 (eRF3)** effective to modulate translation termination. Further, this invention provides an expression vector which comprises a nucleic acid encoding a **MTT1**, a human **Upf1p** protein, a peptidyl eukaryotic **release factor 1 (eRF1)** and a peptidyl eukaryotic **release factor 3 (eRF3)** operably linked to a regulatory element. This invention provides an antibody which binds to the complex comprising an amt. of a **MTT1**, human **Upf1p** protein, a peptidyl eukaryotic **release factor 1 (eRF1)** and a peptidyl eukaryotic **release factor 3 (eRF3)** effective to modulate translation termination. This invention provides an agent which inhibits or modulates the binding of **MTT1** to

eRF3. The agent may inhibit or facilitate the binding of **MTT1** to **eRF3**. Alignment of several RNA **helicases** identifies 9 motifs characteristic of modulators of translation termination.

L4 ANSWER 43 OF 46 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.
on STN DUPLICATE 2

ACCESSION NUMBER: 1998305591 EMBASE
TITLE: A mutated human homologue to yeast **Upf1** protein has a dominant-negative effect on the decay of nonsense-containing mRNAs in mammalian cells.
AUTHOR: Sun X.; Perlick H.A.; Dietz H.C.; Maquat L.E.
CORPORATE SOURCE: L.E. Maquat, Roswell Park Cancer Institute, Department of Genetics, Elm and Carlton Streets, Buffalo, NY 14263, United States. maquat@sc3101.med.buffalo.edu
SOURCE: Proceedings of the National Academy of Sciences of the United States of America, (18 Aug 1998) 95/17 (10009-10014).
Refs: 46
ISSN: 0027-8424 CODEN: PNASA6

COUNTRY: United States
DOCUMENT TYPE: Journal; Article
FILE SEGMENT: 004 Microbiology
LANGUAGE: English
SUMMARY LANGUAGE: English

AB All eukaryotic cells analyzed have developed mechanisms to eliminate the production of mRNAs that prematurely terminate translation. The mechanisms are thought to exist to protect cells from the deleterious effects of in-frame nonsense codons that are generated by routine inefficiencies and inaccuracies in RNA metabolism such as pre-mRNA splicing. Depending on the particular mRNA and how it is produced, nonsense codons can mediate a reduction in mRNA abundance either (i) before its **release** from an association with nuclei into the cytoplasm, presumably but not certainly while the mRNA is being exported to the cytoplasm and translated by cytoplasmic ribosomes, or (ii) in the cytoplasm. Here, we provide evidence for a **factor** that functions to eliminate the production of nonsense-containing RNAs in mammalian cells. The **factor**, variously referred to as Rent1 (regulator of nonsense transcripts) or HUPF1 (human **Upf1** protein), was identified by isolating cDNA for a human homologue to *Saccharomyces cerevisiae Upf1p*, which is a group I RNA **helicase** that functions in the nonsenser mediated decay of mRNA in yeast. Using monkey COS cells and human HeLa cells, we demonstrate that expression of human **Upf1** protein harboring an arginine-to-cysteine mutation at residue 844 within the RNA **helicase** domain acts in a dominant- negative fashion to abrogate the decay of nonsense-containing mRNA that takes place (i) in association with nuclei or (ii) in the cytoplasm. These findings provide evidence that nonsense-mediated mRNA decay is related mechanistically in yeast and in mammalian cells, regardless of the cellular site of decay.

L4 ANSWER 44 OF 46 MEDLINE on STN

ACCESSION NUMBER: 1998283914 MEDLINE
DOCUMENT NUMBER: PubMed ID: 9620853
TITLE: The surveillance complex interacts with the translation **release factors** to enhance termination and degrade aberrant mRNAs.
AUTHOR: Czaplinski K; Ruiz-Echevarria M J; Paushkin S V; Han X; Weng Y; Perlick H A; Dietz H C; Ter-Avanesyan M D; Peltz S W
CORPORATE SOURCE: Department of Molecular Genetics and Microbiology, Robert Wood Johnson Medical School-UMDNJ, USA.
CONTRACT NUMBER: GM48631-01 (NIGMS)
SOURCE: Genes & development, (1998 Jun 1) 12 (11) 1665-77.
Journal code: 8711660. ISSN: 0890-9369.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: Priority Journals
ENTRY MONTH: 199807
ENTRY DATE: Entered STN: 19980713
Last Updated on STN: 19980713

Entered Medline: 19980701

AB The nonsense-mediated mRNA decay pathway is an example of an evolutionarily conserved surveillance pathway that rids the cell of transcripts that contain nonsense mutations. The product of the **UPF1** gene is a necessary component of the putative surveillance complex that recognizes and degrades aberrant mRNAs. Recent results indicate that the **Upf1p** also enhances translation termination at a nonsense codon. The results presented here demonstrate that the yeast and human forms of the **Upf1p** interact with both eukaryotic translation termination factors **eRF1** and **eRF3**. Consistent with **Upf1p** interacting with the **eRFs**, the **Upf1p** is found in the prion-like aggregates that contain **eRF1** and **eRF3** observed in yeast [PSI+] strains. These results suggest that interaction of the **Upf1p** with the peptidyl release factors may be a key event in the assembly of the putative surveillance complex that enhances translation termination, monitors whether termination has occurred prematurely, and promotes degradation of aberrant transcripts.

L4 ANSWER 46 OF 46 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

ACCESSION NUMBER: AAY77814 peptide DGENE

TITLE: New multiprotein complex which can modulate peptidyl transferase activity during translation, useful to treat diseases associated with peptidyl transferase activity e.g. Duchene/Becker Muscular Dystrophy -

INVENTOR: Peltz S; Czaplinski K; Dinman J D

PATENT ASSIGNEE: (UYNE-N)UNIV NEW JERSEY.

PATENT INFO: WO 2000005586 A2 20000203 89p

APPLICATION INFO: WO 1999-US16802 19990722

PRIORITY INFO: US 1998-120435 19980722

DOCUMENT TYPE: Patent

LANGUAGE: English

OTHER SOURCE: 2000-171458 [15]

DESCRIPTION: Yeast **Upf1** protein fragment.

AN AAY77814 peptide DGENE

AB The invention provides a new multiprotein complex which can modulate peptidyl transferase activity during translation. The complex comprises the gene encoding **Helicase B** (HCSB; renamed **MTT1**, for Modulator of Translation Termination) and the conserved proteins known to interact and carry out translation termination in eukaryotic cells, peptidyl eukaryotic release factor (**eRF**) 1 and **eRF3**. The complex can be used to modulate peptidyl transferase activity during translation in a cell. It can be administered therapeutically combined with a carrier in pharmaceutical compositions to treat diseases associated with peptidyl transferase activity, especially diseases resulting from a nonsense or frameshift mutation e.g. beta-thalassemia, beta-globin, Duchene/Becker Muscular Dystrophy etc. It can be used to identify disease conditions involving a defect in the complex, by transfecting cells with encoding nucleic acid and determining the proportion of defective complex before and after transfection. It is also useful to screen for drugs involved in peptidyl transferase activity during translation, inhibiting the interaction between **MTT1** and **eRF3** or involved in enhancing translation termination. Vectors comprising polynucleotides encoding the complex (or antisense sequences) can be constructed and introduced into cells to interfere with complex expression and so modulate the efficiency of translation termination of mRNA and/or degradation of aberrant transcripts in a cell. Agents binding to the complex can be identified and included in therapeutic compositions useful as above, and/or used to modulate peptidyl transferase activity during translation in cells. They are also useful to modulate the efficiency of translation termination of mRNA at a nonsense codon and/or promote degradation of aberrant transcripts in cells. The method can be used to identify agents/ compositions modulating binding to **MTT1**, useful to identify genes. Sequences AAY77813-817 represent protein fragments from yeast superfamily group I **helicases**.

=> dhis

DHIS IS NOT A RECOGNIZED COMMAND

The previous command name entered was not recognized by the system.
For a list of commands available to you in the current file, enter

"HELP COMMANDS" at an arrow prompt (>).

=> d his

(FILE 'HOME' ENTERED AT 19:31:12 ON 28 JUL 2004)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI,
BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA,
CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS,
DDFB, DDFU, DGENE, DRUGB, DRUGMONOG2, ...' ENTERED AT 19:31:26 ON 28 JUL
2004

SEA (HELICAS? OR MTT1?) OR (ERF? OR (RELEAS?(S) FACTO?))

1596 FILE ADISCTI
354 FILE ADISINSIGHT
378 FILE ADISNEWS
6851 FILE AGRICOLA
192 FILE ANABSTR
2954 FILE AQUASCI
1469 FILE BIOBUSINESS
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43311 FILE BIOSIS
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25707 FILE BIOTECHNO
14587 FILE CABA
22758 FILE CANCERLIT
50487 FILE CAPLUS
60171 FILE CEABA-VTB
83 FILE CEN
444 FILE CIN
1196 FILE CONFSCI
306 FILE CROPB
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3955* FILE FEDRIP
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17804 FILE GENBANK
453 FILE HEALSAFE
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72 FILE IMSPRODUCT
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246 FILE KOSMET
20977 FILE LIFESCI
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1131 FILE PROUSSDR
475 FILE RDISCLOSURE

58511 FILE SCISEARCH
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20905 FILE TOXCENTER
43056 FILE USPATFULL
2544 FILE USPAT2
829 FILE VETB
2402 FILE VETU
4638 FILE WPIDS
1925 FILE WPIFV
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944 FILE IPA
171 FILE NAPRALERT
19988 FILE NLDB

L1 QUE (HELICAS? OR MTT1?) OR (ERF? OR (RELEAS?(S) FACTO?))

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PASCAL, BIOSIS, USPATFULL, ESBIOBASE' ENTERED AT 19:35:08 ON 28 JUL 2004
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L3 51 S L2 AND (UPF? OR NAM7? OR SAL1? OR IFS2? OR MOF4? OR NMD2? OR
L4 46 DUP REM L3 (5 DUPLICATES REMOVED)

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|--|------------------|---------------|
| FULL ESTIMATED COST | 60.31 | 63.94 |
| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE ENTRY | TOTAL SESSION |
| CA SUBSCRIBER PRICE | -0.74 | -0.74 |

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STN INTERNATIONAL SESSION SUSPENDED AT 19:44:34 ON 28 JUL 2004


NCBI


Protein

Search for

Limits History Details

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1: NP_542199. UPF3 regulator of...[gi:18375528]

LOCUS NP_542199 483 aa linear PRI 21-DEC-2003
 DEFINITION UPF3 regulator of nonsense transcripts homolog B isoform 1 [Homo sapiens].
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 VERSION NP_542199.1 GI:18375528
 DBSOURCE REFSEQ: accession NM_080632.1
 KEYWORDS .
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (residues 1 to 483)
 REFERENCE
 AUTHORS Gehring,N.H., Neu-Yilik,G., Schell,T., Hentze,M.W. and Kulozik,A.E.
 TITLE Y14 and hUpf3b form an NMD-activating complex
 JOURNAL Mol. Cell 11 (4), 939-949 (2003)
 PUBMED 12718880
 REMARK GeneRIF: A conserved domain of hUpf3b mediates an interaction with the EJC protein Y14. Y14 is required for nonsense-mediated decay induced by tethered hUpf3b.
 2 (residues 1 to 483)
 REFERENCE
 AUTHORS Lykke-Andersen,J., Shu,M.D. and Steitz,J.A.
 TITLE Communication of the position of exon-exon junctions to the mRNA surveillance machinery by the protein RNPS1
 JOURNAL Science 293 (5536), 1836-1839 (2001)
 PUBMED 11546874
 REMARK GeneRIF: binds RNPS1 protein, part of the postslicing complex deposited 5' to exon-exon junctions
 3 (residues 1 to 483)
 REFERENCE
 AUTHORS Kim,V.N., Kataoka,N. and Dreyfuss,G.
 TITLE Role of the nonsense-mediated decay factor hUpf3 in the splicing-dependent exon-exon junction complex
 JOURNAL Science 293 (5536), 1832-1836 (2001)
 PUBMED 11546873
 REMARK GeneRIF: binds to spliced mRNAs upstream of exon-exon junctions; is part of mRNP complexes that are ready for nuclear export and that participate in nonsense-mediated mRNA decay
 4 (residues 1 to 483)
 REFERENCE
 AUTHORS Serin,G., Gersappe,A., Black,J.D., Aronoff,R. and Maquat,L.E.
 TITLE Identification and characterization of human orthologues to Saccharomyces cerevisiae Upf2 protein and Upf3 protein (Caenorhabditis elegans SMG-4)
 JOURNAL Mol. Cell. Biol. 21 (1), 209-223 (2001)
 PUBMED 11113196
 REFERENCE
 AUTHORS Lykke-Andersen,J., Shu,M.D. and Steitz,J.A.
 TITLE Human Upf proteins target an mRNA for nonsense-mediated decay when bound downstream of a termination codon
 JOURNAL Cell 103 (7), 1121-1131 (2000)
 PUBMED 11163187

COMMENT REVIEWED REFSEQ: This record has been curated by NCBI staff. The reference sequence was derived from AF318576.1, AY013251.1 and BI549935.1.

Summary: This gene encodes a protein that is part of a post-splicing multiprotein complex involved in both mRNA nuclear export and mRNA surveillance. The encoded protein is one of two functional homologs to yeast Upf3p. mRNA surveillance detects exported mRNAs with truncated open reading frames and initiates nonsense-mediated mRNA decay (NMD). When translation ends upstream from the last exon-exon junction, this triggers NMD to degrade mRNAs containing premature stop codons. This protein binds to the mRNA and remains bound after nuclear export, acting as a nucleocytoplasmic shuttling protein. It forms with Y14 a complex that binds specifically 20 nt upstream of exon-exon junctions. This gene is located on the long arm of chromosome X. Two splice variants encoding different isoforms have been found for this gene.

Transcript Variant: This variant (1) contains exon 8 and encodes the longer isoform (1), also known as hUpf3-X.

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/db_xref="LocusID:65109"
/db_xref="MIM:300298" |

ORIGIN

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121 dnkggeypai vefapfqkaa kkktkkrdtk vgtiddpey rkflesyatd nekmtstpet
181 lleeieaknr eliakkttpl lsflknkqrm reekreerrr reierkrqre eerrkwkeee
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481 gee

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PubMed

Nucleotide

Protein

Genome

Structure

PMC

Taxonomy

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Details

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20

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File

Get Subsequence

Fe

1: NP_542166. UPF2 regulator of...[gi:18375676]

BLink, Domains, Links

LOCUS NP_542166 1272 aa linear PRI 21-DEC-2003
 DEFINITION UPF2 regulator of nonsense transcripts homolog; regulator of nonsense transcripts 2; yeast Upf2p homolog [Homo sapiens].
 ACCESSION NP_542166
 VERSION NP_542166.1 GI:18375676
 DBSOURCE REFSEQ: accession NM_080599.1
 KEYWORDS .
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RESIDUE 1 (residues 1 to 1272)
 REFERENCE 1 (residues 1 to 1272)
 AUTHORS Serin,G., Gersappe,A., Black,J.D., Aronoff,R. and Maquat,L.E.
 TITLE Identification and characterization of human orthologues to Saccharomyces cerevisiae Upf2 protein and Upf3 protein (Caenorhabditis elegans SMG-4)
 JOURNAL Mol. Cell. Biol. 21 (1), 209-223 (2001)
 PUBMED 11113196
 REFERENCE 2 (residues 1 to 1272)
 AUTHORS Lykke-Andersen,J., Shu,M.D. and Steitz,J.A.
 TITLE Human Upf proteins target an mRNA for nonsense-mediated decay when bound downstream of a termination codon
 JOURNAL Cell 103 (7), 1121-1131 (2000)
 PUBMED 11163187
 REFERENCE 3 (residues 1 to 1272)
 AUTHORS Mendell,J.T., Medghalchi,S.M., Lake,R.G., Noensie,E.N. and Dietz,H.C.
 TITLE Novel Upf2p orthologues suggest a functional link between translation initiation and nonsense surveillance complexes
 JOURNAL Mol. Cell. Biol. 20 (23), 8944-8957 (2000)
 PUBMED 11073994
 COMMENT REVIEWED REFSEQ: This record has been curated by NCBI staff. The reference sequence was derived from AB037829.1 and AW444636.1.

Summary: This gene encodes a protein that is part of a post-splicing multiprotein complex involved in both mRNA nuclear export and mRNA surveillance. mRNA surveillance detects exported mRNAs with truncated open reading frames and initiates nonsense-mediated mRNA decay (NMD). When translation ends upstream from the last exon-exon junction, this triggers NMD to degrade mRNAs containing premature stop codons. This protein is located in the perinuclear area. It interacts with translation release factors and the proteins that are functional homologs of yeast Upf1p and Upf3p. Two splice variants have been found for this gene; both variants encode the same protein.

Transcript Variant: This variant (1) contains a different 5' UTR than variant 2 and is the longer transcript.

FEATURES

Location/Qualifiers

source 1..1272
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="10"
/map="10p14-p13"
Protein 1..1272
/product="UPF2 regulator of nonsense transcripts homolog"
/note="regulator of nonsense transcripts 2; yeast Upf2p homolog"
Region 140..>1024
/region_name="Nonsense-mediated mRNA decay 2 protein [RNA processing and modification]"
/note="KOG2051"
/db_xref="CDD:19837"
Region 168..>320
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/note="MIF4G"
/db_xref="CDD:23388"
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/note="MIF4G"
/db_xref="CDD:23388"
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/region_name="Up-frameshift suppressor 2"
/note="Upf2"
/db_xref="CDD:9609"
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/region_name="Nonsense-mediated mRNA decay 2 protein [RNA processing and modification]"
/note="KOG2051"
/db_xref="CDD:19837"
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/coded_by="NM_080599.1:130..3948"
/note="go_function: RNA binding [goid 0003723] [evidence IEA];
go_process: protein biosynthesis [goid 0006412] [evidence IEA]"
/db_xref="GeneID:26019"
/db_xref="LocusID:26019"
/db_xref="MIM:605529"

ORIGIN

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121 mkekeesiql hqeawerhhl rkelrsknqn apdsrpeenf fsrldsslkk ntafvkkllkt
181 iteqqrdsls hdfnglnlsk yiaeavasiv eaklkisdvn cavhlcslfh qryadfpsl
241 lqvwkkhfea rkeektpnit klrtdlrfia eltivgftd keglsllyeq lkniiinadre
301 shthvsvvis fcrhcggdia glvprkvksa aekfnlsfpp seiispekkqq pfqnllkeyf
361 tsltkhlkrd hrelqnterq nrrilhskge lsedrhkqye efamsyqkll ansqsladll
421 denmpdlpqd kptpeehgpg idiftpgkpg eydleggiwe dedarnfyen lidlkafvp
481 ilfkdnksc qnkesnkddt keakeskenk evsspddlel elenleindd tleleggdea
541 edltkkllde qeqedeeast gshlklivda flqqlpncvn rdlidkaamd fcmmnmntkan
601 rkklvralfi vprqrldllp fyarlvatlh pcmsdvaedl csmlrgdfrf hvrkkdqini
661 etknktvrfi geltkfkkmft knndlhlckm llsdfshhhi emactlletc grflfrspes
721 hlrtsvlleg mmrkkqamhl daryvtmven ayyycnpppa ektvkkkrpp lqeyvrkllly
781 kdlskvtttek vlrqmrklpw qdghevkyvi ccmniwnvk ynsihcvanl laglvlyqed

841 vgihvvdgvl edirlgmevn qpkfnqrris sakflgelyn yrmvesavif rtlysftsf
901 vnpdgspssl dppehlfir lvctildtcg qyfdrgsskr kldcflyfq rywwkksle
961 vwtkdhpfp i didymisdtl ellrpkiklc nsleesirqv qdlereflik lglvndkdsk
1021 dsmtegenle edeeeeegg a eteeqsgnes evneeeeeeg sdnnddegee eeeentdylt
1081 dsnkenetde entevmikgg glkhvpcved edfiqaldkm mlenlqqrsq esvkvhqlkv
1141 aiplhlksql rkgpplggge geaesadttmp fvmltrkgnk qqfkilnvpn ssqlaanhwn
1201 qqaeqeerm rmkkltldin erqeqedyqe mlqslaqrpa pantnrerrp ryqhpkgapn
1261 adlifktggr rr

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NCBI Protein

Entrez PubMed Nucleotide Protein Genome Structure PMC Taxonomy Boo

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BLink, Domains, Links

1: NP_060564. peptide chain rel...[gi:46094014]

LOCUS NP_060564 628 aa linear PRI 12-JUL-2004

DEFINITION peptide chain release factor 3 [Homo sapiens].

ACCESSION NP_060564

VERSION NP_060564.2 GI:46094014

DBSOURCE REFSEQ: accession NM_018094.2

KEYWORDS.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (residues 1 to 628)

AUTHORS Hansen,L.L., Jakobsen,C.G. and Justesen,J.

TITLE Assignment of the human peptide chain release factor 3 (GSPT2) to Xp11.23-->p11.21 and of the distal marker DKS1039 by radiation hybrid mapping

JOURNAL Cytogenet. Cell Genet. 86 (3-4), 250-251 (1999)

PUBMED 10575220

REFERENCE 2 (residues 1 to 628)

AUTHORS Hoshino,S., Imai,M., Mizutani,M., Kikuchi,Y., Hanaoka,F., Ui,M. and Katada,T.

TITLE Molecular cloning of a novel member of the eukaryotic polypeptide chain-releasing factors (eRF). Its identification as eRF3 interacting with eRF1

JOURNAL J. Biol. Chem. 273 (35), 22254-22259 (1998)

PUBMED 9712840

COMMENT VALIDATED REFSEQ: This record has undergone preliminary review of the sequence, but has not yet been subject to final review. The reference sequence was derived from BC036077.1, AJ251548.1, AK001303.1 and AK023155.1.
On Apr 2, 2004 this sequence version replaced gi:8922424.

Summary: GSPT2 is closely related to GSPT1 (MIM 139259), a GTP-binding protein that plays an essential role at the G1- to S-phase transition of the cell cycle in yeast and human cells. GSPT1 is a positive regulator of translational accuracy and, in a binary complex with eRF1 (MIM 600285), functions as a polypeptide chain release factor. [supplied by OMIM].

FEATURES Location/Qualifiers

source 1..628
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/map="Xp11.23-p11.21"

Protein 1..628
/product="peptide chain release factor 3"

Region 201..625
/region_name="Translation elongation factor EF-1alpha (GTPase) [Translation, ribosomal structure and biogenesis]"

Region

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/note="TEF1"
/db_xref="CDD:14378"
201..418
/region_name="Elongation factor Tu GTP binding domain"
/note="GTP_EFTU"
/db_xref="CDD:22868"
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Region

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443..510
/region_name="Elongation factor Tu domain 2"
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/db_xref="CDD:24678"
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Region

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522..624
/region_name="Elongation factor Tu C-terminal domain"
/note="GTP_EFTU_D3"
/db_xref="CDD:23422"
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CDS

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/coded_by="NM_018094.2:197..2083"
/note="go_function: GTP binding [goid 0005525] [evidence IEA];
go_process: protein biosynthesis [goid 0006412] [evidence IEA]"
/db_xref="GeneID:23708"
/db_xref="LocusID:23708"
/db_xref="MIM:300418"
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ORIGIN

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121 legsnsavtm elsepvveng evemaleesw ehskevseae pgggssgdsg ppeesgqemm
181 eekkeirksk svivpsgapk kehvnnvfig hvdagkstig gqimfltgmv dkrtlekyer
241 eakeknretw ylswaldtnq eerdkgktve vgrayfeter khftildapg hksfvpnwmig
301 gasqadlavl visarkgefe tgfekggqtr ehamlaktag vkhlivlink mddptvnwsi
361 eryeeckeekl vpflkkvgfs pkkdihfmpc sgtganike qsdfcpwytg lpfipyldnl
421 pnfnrnsidgp irlpividkyk dmgtvvlgkl esgsifkgqq lvmmmpnkhnv evlgilsddt
481 etdfvapgen lkirlkgiee eeilpgfilc dpsnlchsgsr tfdvqiviie hksiicpgyn
541 avlhihtcie eveitalisl vdkksgeksk trprfvkqdq vciarlrtag ticletfkdf
601 pqmgrftlrd egktiaigkv lklppekd
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//

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The screenshot shows the NCBI Entrez Protein search interface. At the top, there's a navigation bar with links for Entrez, PubMed, Nucleotide, Protein, Genome, Structure, PMC, Taxonomy, and Books. Below the navigation bar is a search bar with the text "Search [Protein] for []". To the right of the search bar are buttons for "Go" and "Clear". Underneath the search bar are buttons for "Display" (set to "default"), "Show" (set to 20), "Send to" (File), "Clipboard", and "Details". A link "Get Subsequence" is also present. On the far right, there are "Links" and "Help" buttons.

Search results for P62495 are displayed. The first result is highlighted with a red border:

1: P62495. Eukaryotic peptid...[gi:50402099]

LOCUS P62495 437 aa linear PRI 01-OCT-2004
DEFINITION Eukaryotic peptide chain release factor subunit 1 (eRF1)
 (Eukaryotic release factor 1) (TB3-1) (C11 protein).
ACCESSION P62495
VERSION P62495 GI:50402099
DBSOURCE swissprot: locus ERF1_HUMAN, accession P62495;
 class: standard.
 extra accessions:P46055,created: Nov 1, 1995.
 sequence updated: Nov 1, 1995.
 annotation updated: Oct 1, 2004.
 xrefs: gi: [338686](#), gi: [338687](#), gi: [1491703](#), gi: [1491704](#), gi: [1890299](#), gi: [1890300](#), gi: [5499720](#), gi: [5499721](#), gi: [1082824](#), pdb accession 1DT9
 xrefs (non-sequence databases): GenewHGNC:3477, MIM [600285](#), GO0005737, GO0003723, GO0003747, GO0006449, InterProIPR004403, InterProIPR005140, InterProIPR005141, InterProIPR005142, PfamPF03463, PfamPF03464, PfamPF03465, TIGRFAMsTIGR00108
KEYWORDS 3D-structure; Protein biosynthesis.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (residues 1 to 437)
AUTHORS Grenett,H.E., Bounelis,P. and Fuller,G.M.
TITLE Identification of a human cDNA with high homology to yeast omnipotent suppressor 45
JOURNAL Gene 110 (2), 239-243 (1992)
MEDLINE [92165066](#)
PUBMED [1537561](#)
REMARK SEQUENCE FROM N.A.
REFERENCE 2 (residues 1 to 437)
AUTHORS Frolova,L., Le Goff,X., Rasmussen,H.H., Cheperegin,S., Drugeon,G., Kress,M., Arman,I., Haenni,A.L., Celis,J.E., Philippe,M. et al.
TITLE A highly conserved eukaryotic protein family possessing properties of polypeptide chain release factor
JOURNAL Nature 372 (6507), 701-703 (1994)
MEDLINE [95082951](#)
PUBMED [7990965](#)
REMARK REVISIONS, AND FUNCTION.
REFERENCE 3 (residues 1 to 437)
AUTHORS Andjelkovic,N., Zolnierowicz,S., Van Hoof,C., Goris,J. and Hemmings,B.A.
TITLE The catalytic subunit of protein phosphatase 2A associates with the translation termination factor eRF1
JOURNAL EMBO J. 15 (24), 7156-7167 (1996)
MEDLINE [97157506](#)
PUBMED [9003791](#)
REMARK SEQUENCE FROM N.A.
 TISSUE=Brain

REFERENCE 4 (residues 1 to 437)
AUTHORS Guenet,L., Toutain,B., Guilleret,I., Chauvel,B., Deaven,L.L.,
Longmire,J.L., Le Gall,J.Y., David,V. and Le Treut,A.
TITLE Human release factor eRF1: structural organisation of the unique
functional gene on chromosome 5 and of the three processed
pseudogenes
JOURNAL FEBS Lett. 454 (1-2), 131-136 (1999)
MEDLINE 99339455
PUBMED 10413110
REMARK SEQUENCE FROM N.A.
REFERENCE 5 (residues 1 to 437)
AUTHORS Song,H., Mugnier,P., Das,A.K., Webb,H.M., Evans,D.R., Tuite,M.F.,
Hemmings,B.A. and Barford,D.
TITLE The crystal structure of human eukaryotic release factor
eRF1--mechanism of stop codon recognition and peptidyl-tRNA
hydrolysis
JOURNAL Cell 100 (3), 311-321 (2000)
MEDLINE 20139983
PUBMED 10676813
REMARK X-RAY CRYSTALLOGRAPHY (2.7 ANGSTROMS).
COMMENT On Jul 20, 2004 this sequence version replaced gi:1169547.

This SWISS-PROT entry is copyright. It is produced through a
collaboration between the Swiss Institute of Bioinformatics and
the EMBL outstation - the European Bioinformatics Institute.
The original entry is available from <http://www.expasy.ch/sprot>
and <http://www.ebi.ac.uk/sprot>

[FUNCTION] Directs the termination of nascent peptide synthesis
(translation) in response to the termination codons UAA, UAG and
UGA.

[SUBUNIT] Heterodimer of two subunits, one of which binds GTP.

[SUBCELLULAR LOCATION] Cytoplasmic.

[SIMILARITY] Belongs to the eukaryotic release factor 1 family.

FEATURES Location/Qualifiers
source 1..437
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/db_xref="taxon:9606"
gene 1..437
/gene="Name=ETF1"
/note="synonyms: Synonyms=ERF1,, RF1"
Protein 1..437
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Region 7..9
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Region 26
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Region 34..39
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Region 45..59
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Region 60..61
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Region 83..84
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Region 107..114
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Region 134..139
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Region 145..150
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Region 158..162
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Region 163..164
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Region 188..197
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Region 198..203
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Region 319..323
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Region 374..380
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Region 383..386
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Region 405..408
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Region 409..412
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/region_name="Beta-strand region"

ORIGIN
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121 ntstlylcndnk fhteaaltall sddskfgfiv idgsgalfgt lqgnntrevlh kftvdlpkkh
181 grggqsalrf arlrmekrhn yvrkvaetav qlfisgdkvn vaglqlagsa dfktelsqsd
241 mfdqrlqskv lklvdisygg engfnqaiel stevlsvnkvf iqekkligry fdeisqdtgk
301 ycfgvedtlk alemgaveil ivyenldimr yvlhcqgtee ekilyltpeq ekdkshftdk
361 etggehelie smpllewfyan nykkfgatle ivtdksqegs qfvkgfggig gilryrvdfq
421 gmeeyqggdde fflddy
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